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INVESTOR IN PEOPLE

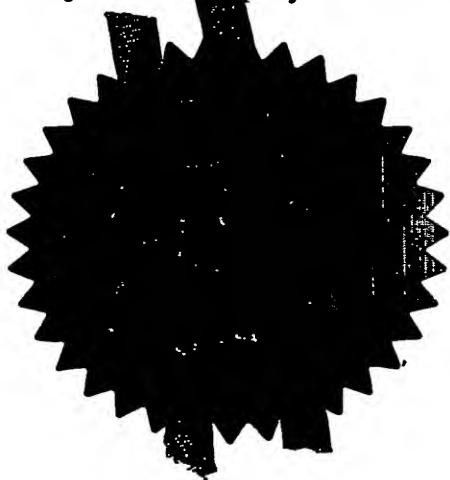
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R. McChesney

13 June 2003



16 JUL 02 E73376-1 002934
PO 0216321.0

Request for grant of a patent

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The Patent Office
Cardiff Road
Newport
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1. Your reference 100771

2. Patent application number
(The Patent Office will fill in this part) 0216321.0

13 JUL 2002

3. Full name, address and postcode of the or of each applicant *(underline all surnames)*
AstraZeneca AB
S-151 85 Sodertalje
Sweden

Patents ADP number *(if you know it)*

If the applicant is a corporate body, give the country/state of its incorporation

Sweden

79 570 21 000

4. Title of the invention

THERAPEUTIC TREATMENT

5. Name of your agent *(if you have one)*

Lucy Clare Padgett

"Address for service" in the United Kingdom to which all correspondence should be sent
(including the postcode)

AstraZeneca UK Limited
Global Intellectual Property
Mereside, Alderley Park
Macclesfield
Cheshire SK10 4TG

Patents ADP number *(if you know it)*

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and *(if you know it)* the or each application number

Country	Priority application number <i>(if you know it)</i>	Date of filing <i>(day / month / year)</i>
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7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application	Date of filing <i>(day / month / year)</i>
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8. Is a statement of inventorship and of right to grant of a patent required in support of this request? *(Answer 'Yes' if:*

- a) *any applicant named in part 3 is not an inventor, or*
- b) *there is an inventor who is not named as an applicant, or*
- c) *any named applicant is a corporate body.*

See note (d))

Patents Form 1/77

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Continuation sheets of this form

Description

36

Claim(s)

03

Abstract

01

Drawing(s)

10. If you are also filing any of the following, state how many against each item.

Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (*Patents Form 7/77*)

Request for preliminary examination and search (*Patents Form 9/77*)

Request for substantive examination
(*Patents Form 10/77*)

Any other documents
(please specify)

11.

I/We request the grant of a patent on the basis of this application.

Signature

Authorised Signatory

Date

12/07/2002

12. Name and daytime telephone number of person to contact in the United Kingdom

Jennifer C Bennett - 01625 230148

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THEAPEUTIC TREATMENT

The present invention relates to combination treatments comprising a metal salt and compounds that inhibit ileal bile acid transport (IBAT) inhibitory activity wherein the metal
5 salt is formulated to release in the terminal ileum, caecum and/or the colon. These combination treatments are useful in preventing diarrhoea that would result from excess bile acids in the intestine following administration of an effective amount an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a warm-blooded animal, such as man. The invention also relates to pharmaceutical
10 compositions containing these combinations and to their use in the manufacture of medicaments. These combinations have value in the treatment of disease states associated with hyperlipidaemic conditions.

It is well-known that hyperlipidaemic conditions associated with elevated concentrations of total cholesterol and LDL cholesterol are major risk factors for
15 cardiovascular atherosclerotic disease (Circulation 1999, 100, 1930-1938 and Circulation, 1999, 100, 1134-46). To reduce the risk and the total mortality due to cardiovascular disease, the reduction of plasma lipids, particularly LDL cholesterol, is now recognized as an important therapeutic goal (N Engl J Med. 1995; 332:5, 12-21).

Interfering with the circulation of bile acids within the lumen of the intestinal tracts
20 has also been found to reduce the level of cholesterol. Bile acids are synthesized in the liver from cholesterol and secreted into the bile. They are actively recycled (>95%) from the small intestine back to the liver. Previous established therapies have involved, for example, treatment with bile acid binders, such as resins. Frequently used bile acid binders are for instance cholestyramine and cholestipol.

25 Another proposed therapy (Current Opinion on Lipidology, 1999, 10, 269-74) involves the treatment with substances with an IBAT inhibitory effect. Theoretically, IBAT inhibitors should have similar therapeutic effect as the resins but they might also be expected to have attractive advantages. First, it should be possible to administer IBAT inhibitors as tablets at the same dose intervals as statins. Second, a direct inhibition of the transport of bile
30 acids across the ileum should be advantageous in situations when IBAT is upregulated. However available data on the effects of IBAT inhibitors is limited. Several IBAT agents have previously been shown to promote the faecal excretion of bile acids and to reduce plasma cholesterol. The proposed mechanism for the hypolipidaemic action of these

compounds is by an induced number of hepatic LDL receptors due to the increased consumption of hepatic cholesterol caused by a compensatory increased bile acid synthesis (Arterioscler Thromb Vasc Biol. 1998; 18: 1304-11).

However, bile acids that are not recycled in the intestines induce irritation of the 5 intestinal luminal surfaces, at least at higher concentrations. This is seen for example in chronic diarrhoea, and in post infectious diarrhoea with deficient uptake of bile acids, after continuous bile acid secretion following cholecystectomy and after resection of the distal ileum. *In vivo* dosing of IBAT compounds may give rise to these side effects either in certain patients or at high enough doses, i.e. irritation of the intestine would be induced, resulting in 10 diarrhoea. The present invention ameliorates this problem.

Furthermore, if chronic diarrhoea was a side effect, then it is possible that these 15 compounds would not be suitable for administering to patients at all (or at least at high enough doses to give a therapeutic effect), despite their efficacy. The present invention therefore provides the additional advantage that it opens up treatment with an IBAT inhibitor to a particular patient population where it might otherwise have not been possible to use these compounds.

Patients suffering from bile acid induced diarrhoea caused by intestinal bypass for 20 example have previously been treated with large doses (2-4 g) of a calcium salt (Reference: Steinbach et al Eur. J of Gastroenterology & Hepatology 1996, 8:559-562). A 2-4 g dose of a salt is too large for convenient dosing regimen, and patient compliance with this regime would be in doubt. This dose is also too large to make a single tablet made up of the IBAT 25 inhibitor and the salt, which is one aspect of the present invention. A formulation which delivers the metal salt with a targeted release to the terminal ileum, caecum and/or the colon would allow a much lower dose of the salt to be used because there will be no loss of the metal salt due to absorption or binding to other components in the small intestine. Therefore it 30 should be possible to formulate a convenient combination regimen, either a single combination tablet or otherwise.

In the literature IBAT inhibitors are often referred to by different names. It is to be understood that where IBAT inhibitors are referred to herein, this term also encompasses 30 compounds known in the literature as:

- i) ileal apical sodium co-dependent bile acid transporter (ASBT) inhibitors;
- ii) bile acid transporter (BAT) inhibitors;
- iii) ileal sodium/bile acid cotransporter system inhibitors;

- iv) apical sodium-bile acid cotransporter inhibitors;
 - v) ileal sodium-dependent bile acid transport inhibitors;
 - vi) bile acid reabsorption (BARI's) inhibitors; and
 - vii) sodium bile acid transporter (SBAT) inhibitors;
- 5 where they act by inhibition of IBAT.

Accordingly the present invention provides a combination which comprises an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon.

10 Herein, where the term "combination" is used it is to be understood that this refers to simultaneous, separate or sequential administration. In one aspect of the invention "combination" refers to simultaneous administration. In another aspect of the invention "combination" refers to separate administration. In a further aspect of the invention "combination" refers to sequential administration. Where the administration is sequential or 15 separate, the delay in administering the second component should not be such as to lose the benefit of the combination.

The combination of the present invention may either be in the form of a fixed combination with the IBAT inhibitor, in which case both the IBAT inhibitor and the metal salt are formulated to release in the colon, or a free combination wherein only the metal salt is 20 formulated to release in the colon.

Suitable metals in the metal salt include any pharmaceutically acceptable multivalent metal ion. In one aspect of the invention these metals are calcium, aluminium, iron, copper, zinc, magnesium, manganese or tin salts. In another aspect of the invention these metals are Ca(II), Al(III), Fe(II), Fe(III), Cu(II), Zn(II), Mg(II), Mn(II) or Sn(II) salts. In a further aspect 25 of the invention the metal in the metal salt is calcium. In another aspect the metal in the metal salt is Ca(II). The salt may be any suitable pharmaceutically acceptable salt. In one aspect the salt is acetate, ascorbate, carbonate, chloride, citrate, gluconate, lactate, nitrate, oxalate, phosphate or sulphate. Suitable metal salts include calcium phosphate, calcium lactate, calcium carbonate, calcium gluconate and calcium acetate, particularly calcium phosphate.

30 It is to be understood that the combination of the present invention includes the situation where there is one metal salt in the combination with the IBAT inhibitor. In addition the combination of the present invention includes the situation where there are one or more metal salts in the combination with the IBAT inhibitor. In this case the salts may be one or

more different salts of the same metal, one or more of the same salt of different metals or one or more different salts of different metals.

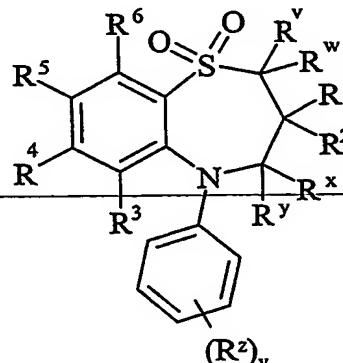
Suitable compounds possessing IBAT inhibitory activity have been described, see for instance the compounds described in WO 93/16055, WO 94/18183, WO 94/18184, WO 5 96/05188, WO 96/08484, WO 96/16051, WO 97/33882, WO 98/38182, WO 99/35135, WO 98/40375, WO 99/35153, WO 99/64409, WO 99/64410, WO 00/01687, WO 00/47568, WO 00/61568, WO 01/68906, DE 19825804, WO 00/38725, WO 00/38726, WO 00/38727, WO 00/38728, WO 00/38729, WO 01/68906, WO 01/66533 and EP 864 582 and the contents of these patent applications are incorporated herein by reference.

10 Particular classes of IBAT inhibitors suitable for use in the present invention are benzothiepines, and the compounds described in the claims, particularly claim 1, of WO 00/01687, WO 96/08484 and WO 97/33882 are incorporated herein by reference. Other suitable classes of IBAT inhibitors are the 1,2-benzothiazepines, 1,4-benzothiazepines and 1,5-benzothiazepines. A further suitable class of IBAT inhibitors is the 1,2,5-benzothiadiazepines.

15

One particular suitable compound possessing IBAT inhibitory activity is (3*R*,5*R*)-3-butyl-3-ethyl-1,1-dioxido-5-phenyl-2,3,4,5-tetrahydro-1,4-benzothiazepin-8-yl β-D-glucopyranosiduronic acid (EP 864 582).

20 Additional suitable compounds possessing IBAT inhibitory activity have the following structure of formula (AI):



(AI)

wherein:

R^v and R^w are independently selected from hydrogen or C₁₋₆alkyl;

25 R^1 and R^2 are independently selected from C₁₋₆alkyl;

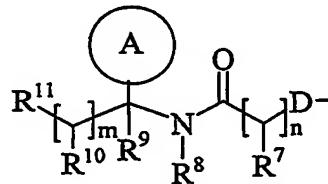
R^x and R^y are independently selected from hydrogen or C₁₋₆alkyl, or one of R^x and R^y is hydrogen or C₁₋₆alkyl and the other is hydroxy or C₁₋₆alkoxy;

\mathbf{R}^2 is selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{1-6} alkoxy, C_{1-6} alkanoyl, C_{1-6} alkanoyloxy, $N-(C_{1-6}$ alkyl)amino, $N,N-(C_{1-6}$ alkyl)₂amino, C_{1-6} alkanoylamino, $N-(C_{1-6}$ alkyl)carbamoyl, $N,N-(C_{1-6}$ alkyl)₂carbamoyl, C_{1-6} alkylS(O)_a wherein a is 0 to 2, C_{1-6} alkoxycarbonyl,

- 5 C_{1-6} alkoxycarbonylamino, ureido, $N'-(C_{1-6}$ alkyl)ureido, $N-(C_{1-6}$ alkyl)ureido, $N',N'-(C_{1-6}$ alkyl)₂ureido, $N'-(C_{1-6}$ alkyl)- $N-(C_{1-6}$ alkyl)ureido, $N',N'-(C_{1-6}$ alkyl)₂- $N-(C_{1-6}$ alkyl)ureido, $N-(C_{1-6}$ alkyl)sulphamoyl and $N,N-(C_{1-6}$ alkyl)₂sulphamoyl;

v is 0-5;

- 10 one of \mathbf{R}^4 and \mathbf{R}^5 is a group of formula (AIA):



(AIA)

- \mathbf{R}^3 and \mathbf{R}^6 and the other of \mathbf{R}^4 and \mathbf{R}^5 are independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, C_{1-4} alkanoyl, C_{1-4} alkanoyloxy, $N-(C_{1-4}$ alkyl)amino, $N,N-(C_{1-4}$ alkyl)₂amino, C_{1-4} alkanoylamino, $N-(C_{1-4}$ alkyl)carbamoyl, $N,N-(C_{1-4}$ alkyl)₂carbamoyl, C_{1-4} alkylS(O)_a wherein a is 0 to 2, C_{1-4} alkoxycarbonyl, $N-(C_{1-4}$ alkyl)sulphamoyl and $N,N-(C_{1-4}$ alkyl)₂sulphamoyl; wherein \mathbf{R}^3 and \mathbf{R}^6 and the other of \mathbf{R}^4 and \mathbf{R}^5 may be optionally substituted on carbon by one or more \mathbf{R}^{16} ;

- 20 D is $-O-$, $-N(R^a)-$, $-S(O)_b-$ or $-CH(R^a)-$; wherein R^a is hydrogen or C_{1-6} alkyl and b is 0-2;

Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from \mathbf{R}^{17} ;

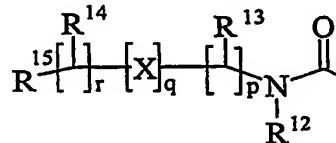
- 25 \mathbf{R}^7 is hydrogen, C_{1-4} alkyl, carbocyclyl or heterocyclyl; wherein \mathbf{R}^7 is optionally substituted by one or more substituents selected from \mathbf{R}^{18} ;

\mathbf{R}^8 is hydrogen or C_{1-4} alkyl;

\mathbf{R}^9 is hydrogen or C_{1-4} alkyl;

\mathbf{R}^{10} is hydrogen, C_{1-4} alkyl, carbocyclyl or heterocyclyl; wherein \mathbf{R}^{10} is optionally substituted by one or more substituents selected from \mathbf{R}^{19} ;

R¹¹ is carboxy, sulpho, sulphino, phosphono, tetrazolyl, -P(O)(OR^c)(OR^d), -P(O)(OH)(OR^c), -P(O)(OH)(R^d) or -P(O)(OR^c)(R^d) wherein R^c and R^d are independently selected from C₁₋₆alkyl; or R¹¹ is a group of formula (AIB):



5

(AIB)

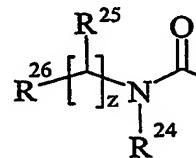
wherein:

X is -N(R^a)-, -N(R^a)C(O)-, -O-, and -S(O)_a-; wherein a is 0-2 and R^a is hydrogen or C₁₋₄alkyl;

R¹² is hydrogen or C₁₋₄alkyl;

10 **R¹³** and **R¹⁴** are independently selected from hydrogen, C₁₋₄alkyl, carbocyclyl, heterocyclyl or R²³; wherein said C₁₋₄alkyl, carbocyclyl or heterocyclyl may be independently optionally substituted by one or more substituents selected from R²⁰;

R¹⁵ is carboxy, sulpho, sulphino, phosphono, tetrazolyl, -P(O)(OR^e)(OR^f), -P(O)(OH)(OR^e), -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from C₁₋₆alkyl; or R¹⁵ is a group of formula (AIC):



(AIC)

wherein:

R²⁴ is selected from hydrogen or C₁₋₄alkyl;

20 **R²⁵** is selected from hydrogen, C₁₋₄alkyl, carbocyclyl, heterocyclyl or R²⁷; wherein said C₁₋₄alkyl, carbocyclyl or heterocyclyl may be independently optionally substituted by one or more substituents selected from R²⁸;

R²⁶ is selected from carboxy, sulpho, sulphino, phosphono, tetrazolyl, -P(O)(OR^g)(OR^h), -P(O)(OH)(OR^g), -P(O)(OH)(R^g) or -P(O)(OR^g)(R^h) wherein R^g and R^h are independently selected from C₁₋₆alkyl;

p is 1-3; wherein the values of R¹³ may be the same or different;

q is 0-1;

r is 0-3; wherein the values of R¹⁴ may be the same or different;

m is 0-2; wherein the values of **R**¹⁰ may be the same or different;

n is 1-3; wherein the values of **R**⁷ may be the same or different;

z is 0-3; wherein the values of **R**²⁵ may be the same or different;

R¹⁶, **R**¹⁷ and **R**¹⁸ are independently selected from halo, nitro, cyano, hydroxy, amino,

- 5 carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino, C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a wherein **a** is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl and N,N-(C₁₋₄alkyl)₂sulphamoyl; wherein **R**¹⁶, **R**¹⁷ and **R**¹⁸ may be independently optionally substituted on carbon by one or more **R**²¹;

10 **R**¹⁹, **R**²⁰, **R**²³, **R**²⁷ and **R**²⁸ are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino, C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a

- 15 wherein **a** is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl, N,N-(C₁₋₄alkyl)₂sulphamoyl, carbocyclyl, heterocyclyl, sulpho, sulphino, amidino, phosphono, -P(O)(OR^a)(OR^b), -P(O)(OH)(OR^a), -P(O)(OH)(R^a) or -P(O)(OR^a)(R^b), wherein **R**^a and **R**^b are independently selected from C₁₋₆alkyl; wherein **R**¹⁹, **R**²⁰, **R**²³, **R**²⁷ and **R**²⁸ may be independently optionally substituted on carbon by one or more **R**²²;

- 20 **R**²¹ and **R**²² are independently selected from halo, hydroxy, cyano, carbamoyl, ureido, amino, nitro, carboxy, carbamoyl, mercapto, sulphamoyl, trifluoromethyl, trifluoromethoxy, methyl, ethyl, methoxy, ethoxy, vinyl, allyl, ethynyl, methoxycarbonyl, formyl, acetyl, formamido, acetylamino, acetoxy, methylamino, dimethylamino, N-methylcarbamoyl, N,N-dimethylcarbamoyl, methylthio, methylsulphinyl, mesyl, N-methylsulphamoyl and N,N-dimethylsulphamoyl;

25 or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Particular compounds of formula (A1) are:

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(carboxymethyl) carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

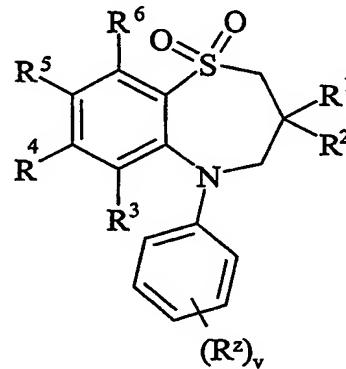
- 30 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-α-[N'-(carboxymethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(2-sulphoethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

- 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(*N*-{(R)-1'-phenyl-1'-[*N'*-(2-sulphoethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[*N'*-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 5 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[*N'*-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[*N'*-(2-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[*N'*-(2-carboxyethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 10 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[*N'*-(5-carboxypentyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[*N'*-(2-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 15 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(*N*-{ α -[*N'*-(2-sulphoethyl)carbamoyl]-2-fluorobenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[*N'*-(R)-(2-hydroxy-1-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[*N'*-(R)-(2-hydroxy-1-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 20 20 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(*N*-[(R)- α -{*N'*-{(R)-1-[*N''*-(R)-(2-hydroxy-1-carboxyethyl)carbamoyl]-2-hydroxyethyl}carbamoyl}benzyl]carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
-
- 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(*N*-{ α -[*N'*-(carboxymethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 25 25 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(*N*-{ α -[*N'*-((ethoxy)(methyl)phosphorylmethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-{*N*-[(R)- α -{*N'*-{2-[hydroxy(methyl)phosphoryl]ethyl}carbamoyl}benzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 30 30 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[*N'*-(2-methylthio-1-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{N-[*(R)*- α -(*N'*-{2-[(methyl)(ethyl)phosphoryl]ethyl}carbamoyl)-4-hydroxybenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 5 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{N-[*(R)*- α -(*N'*-{2-[(methyl)(hydroxy)phosphoryl]ethyl}carbamoyl)-4-hydroxybenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(*N*-{(R)- α -[(R)-*N'*-(2-methylsulphinyl-1-carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine; and
- 10 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methoxy-8-[*N*-{(R)- α -[*N'*-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine; or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Further suitable compounds possessing IBAT inhibitory activity have the following structure of formula (BI):



15

(BI)

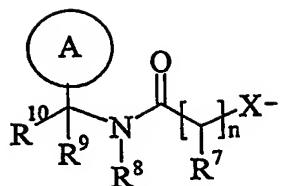
wherein:

One of R¹ and R² are selected from hydrogen or C₁₋₆alkyl and the other is selected from C₁₋₆alkyl;

- 20 R² is selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₁₋₆alkoxy, C₁₋₆alkanoyl, C₁₋₆alkanoyloxy, N-(C₁₋₆alkyl)amino, N,N-(C₁₋₆alkyl)₂amino, C₁₋₆alkanoylamino, N-(C₁₋₆alkyl)carbamoyl, N,N-(C₁₋₆alkyl)₂carbamoyl, C₁₋₆alkylS(O)_a wherein a is 0 to 2, C₁₋₆alkoxycarbonyl, N-(C₁₋₆alkyl)sulphamoyl and N,N-(C₁₋₆alkyl)₂sulphamoyl;

25 v is 0-5;

one of R⁴ and R⁵ is a group of formula (BIA):



(BIA)

R^3 and R^6 and the other of R^4 and R^5 are independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-6} alkyl,

- 5 C_{2-6} alkenyl, C_{2-6} alkynyl, C_{1-6} alkoxy, C_{1-6} alkanoyl, C_{1-6} alkanoyloxy, $N-(C_{1-6}$ alkyl)amino, $N,N-(C_{1-6}$ alkyl)₂amino, C_{1-6} alkanoylamino, $N-(C_{1-6}$ alkyl)carbamoyl, $N,N-(C_{1-6}$ alkyl)₂carbamoyl, C_{1-6} alkylS(O)_a wherein a is 0 to 2, C_{1-6} alkoxycarbonyl, $N-(C_{1-6}$ alkyl)sulphamoyl and $N,N-(C_{1-6}$ alkyl)₂sulphamoyl; wherein R^3 and R^6 and the other of R^4 and R^5 may be optionally substituted on carbon by one or more R^{17} ;
- 10 X is $-O-$, $-N(R^a)-$, $-S(O)_b-$ or $-CH(R^a)-$; wherein R^a is hydrogen or C_{1-6} alkyl and b is 0-2;

Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted on carbon by one or more substituents selected from R^{18} ;

- 15 R^7 is hydrogen, C_{1-6} alkyl, carbocyclyl or heterocyclyl; wherein R^7 is optionally substituted on carbon by one or more substituents selected from R^{19} ; and wherein if said heterocyclyl contains an -NH- group, that nitrogen may be optionally substituted by a group selected from R^{20} ;

R^8 is hydrogen or C_{1-6} alkyl;

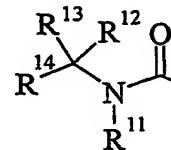
R^9 is hydrogen or C_{1-6} alkyl;

- 20 R^{10} is hydrogen, halo, nitro, cyano, hydroxy, amino, carbamoyl, mercapto, sulphamoyl, hydroxyaminocarbonyl, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, C_{1-10} alkoxy,

C_{1-10} alkanoyl, C_{1-10} alkanoyloxy, $N-(C_{1-10}$ alkyl)amino, $N,N-(C_{1-10}$ alkyl)₂amino, $N,N,N-(C_{1-10}$ alkyl)₃ammonio, C_{1-10} alkanoylamino, $N-(C_{1-10}$ alkyl)carbamoyl, $N,N-(C_{1-10}$ alkyl)₂carbamoyl, C_{1-10} alkylS(O)_a wherein a is 0 to 2, $N-(C_{1-10}$ alkyl)sulphamoyl,

- 25 $N,N-(C_{1-10}$ alkyl)₂sulphamoyl, $N-(C_{1-10}$ alkyl)sulphamoylamino, $N,N-(C_{1-10}$ alkyl)₂sulphamoylamino, C_{1-10} alkoxycarbonylamino, carbocyclyl, carbocyclyl C_{1-10} alkyl, heterocyclyl, heterocyclyl C_{1-10} alkyl, carbocyclyl-(C_{1-10} alkylene)_p- R^{21} -(C_{1-10} alkylene)_q- or heterocyclyl-(C_{1-10} alkylene)_r- R^{22} -(C_{1-10} alkylene)_s-; wherein R^{10} is optionally substituted on carbon by one or more substituents selected from R^{23} ; and wherein if said heterocyclyl
- 30

contains an -NH- group, that nitrogen may be optionally substituted by a group selected from R²⁴; or R¹⁰ is a group of formula (BIB):



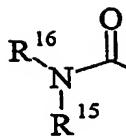
(BIB)

5 wherein:

R¹¹ is hydrogen or C₁₋₆alkyl;

- 10 R¹² and R¹³ are independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carbamoyl, mercapto, sulphamoyl, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₁₋₁₀alkoxy, C₁₋₁₀alkanoyl, C₁₋₁₀alkanoyloxy, N-(C₁₋₁₀alkyl)amino, N,N-(C₁₋₁₀alkyl)₂amino, C₁₋₁₀alkanoylamino, N-(C₁₋₁₀alkyl)carbamoyl, N,N-(C₁₋₁₀alkyl)₂carbamoyl, C₁₋₁₀alkylS(O)_a wherein a is 0 to 2, N-(C₁₋₁₀alkyl)sulphamoyl, N,N-(C₁₋₁₀alkyl)₂sulphamoyl, N-(C₁₋₁₀alkyl)sulphamoylamino, N,N-(C₁₋₁₀alkyl)₂sulphamoylamino, carbocyclyl or heterocyclyl; wherein R¹² and R¹³ may be independently optionally substituted on carbon by one or more substituents selected from R²⁵; and wherein if said heterocyclyl contains an -NH- group, that nitrogen may be optionally substituted by a group selected from R²⁶;

- 15 R¹⁴ is selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carbamoyl, mercapto, sulphamoyl, hydroxyaminocarbonyl, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₁₋₁₀alkoxy, C₁₋₁₀alkanoyl, C₁₋₁₀alkanoyloxy, N-(C₁₋₁₀alkyl)amino, N,N-(C₁₋₁₀alkyl)₂amino, N,N,N-(C₁₋₁₀alkyl)₃ammonio, C₁₋₁₀alkanoylamino, N-(C₁₋₁₀alkyl)carbamoyl, N,N-(C₁₋₁₀alkyl)₂carbamoyl, C₁₋₁₀alkylS(O)_a wherein a is 0 to 2, N-(C₁₋₁₀alkyl)sulphamoyl, N,N-(C₁₋₁₀alkyl)₂sulphamoyl, N-(C₁₋₁₀alkyl)sulphamoylamino, N,N-(C₁₋₁₀alkyl)₂sulphamoylamino, C₁₋₁₀alkoxycarbonylamino, carbocyclyl, carbocyclylC₁₋₁₀alkyl, heterocyclyl, heterocyclylC₁₋₁₀alkyl, carbocyclyl-(C₁₋₁₀alkylene)_b-R²⁷-(C₁₋₁₀alkylene)_q- or 20 heterocyclyl-(C₁₋₁₀alkylene)_r-R²⁸-(C₁₋₁₀alkylene)_s-; wherein R¹⁴ may be optionally substituted on carbon by one or more substituents selected from R²⁹; and wherein if said heterocyclyl contains an -NH- group, that nitrogen may be optionally substituted by a group selected from R³⁰; or R¹⁴ is a group of formula (BIC):



(BIC)

R¹⁵ is hydrogen or C₁₋₆alkyl;

5 R¹⁶ is hydrogen or C₁₋₆alkyl; wherein R¹⁶ may be optionally substituted on carbon by one or more groups selected from R³¹;

n is 1-3; wherein the values of R⁷ may be the same or different;

10 R¹⁷, R¹⁸, R¹⁹, R²³, R²⁵, R²⁹ or R³¹ are independently selected from halo, nitro, cyano, hydroxy, amino, carbamoyl, mercapto, sulphamoyl, hydroxyaminocarbonyl, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₁₋₁₀alkoxy, C₁₋₁₀alkanoyl, C₁₋₁₀alkanoyloxy, N-(C₁₋₁₀alkyl)amino, N,N-(C₁₋₁₀alkyl)₂amino, N,N,N-(C₁₋₁₀alkyl)₃ammonio, C₁₋₁₀alkanoylamino, N-(C₁₋₁₀alkyl)carbamoyl, N,N-(C₁₋₁₀alkyl)₂carbamoyl, C₁₋₁₀alkylS(O)_a wherein a is 0 to 2, N-(C₁₋₁₀alkyl)sulphamoyl, N,N-(C₁₋₁₀alkyl)₂sulphamoyl, N-(C₁₋₁₀alkyl)sulphamoylamino, N,N-(C₁₋₁₀alkyl)₂sulphamoylamino, C₁₋₁₀alkoxycarbonylamino, carbocyclyl, carbocyclylC₁₋₁₀alkyl, heterocyclyl, heterocyclylC₁₋₁₀alkyl,

15 carbocyclyl-(C₁₋₁₀alkylene)_p-R³²-(C₁₋₁₀alkylene)_q- or heterocyclyl-(C₁₋₁₀alkylene)_r-R³³-(C₁₋₁₀alkylene)_s; wherein R¹⁷, R¹⁸, R¹⁹, R²³, R²⁵, R²⁹ or R³¹ may be independently optionally substituted on carbon by one or more R³⁴; and wherein if said heterocyclyl contains an -NH- group, that nitrogen may be optionally substituted by a group selected from R³⁵;

20 R²¹, R²², R²⁷, R²⁸, R³² or R³³ are independently selected from -O-, -NR³⁶-, -S(O)_x-, -NR³⁶C(O)NR³⁶-, -NR³⁶C(S)NR³⁶-, -OC(O)N=C-, -NR³⁶C(O)- or -C(O)NR³⁶-, wherein R³⁶ is selected from hydrogen or C₁₋₆alkyl, and x is 0-2;

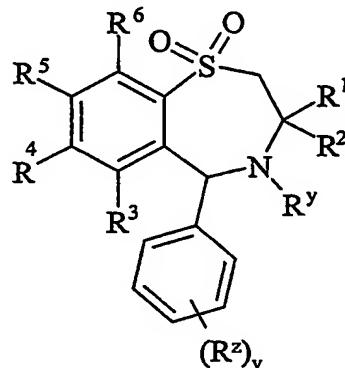
p, q, r and s are independently selected from 0-2;

R³⁴ is selected from halo, hydroxy, cyano, carbamoyl, ureido, amino, nitro, carbamoyl,

25 mercapto, sulphamoyl, trifluoromethyl, trifluoromethoxy, methyl, ethyl, methoxy, ethoxy, vinyl, allyl, ethynyl, formyl, acetyl, formamido, acetylarnino, acetoxy, methylarnino, dimethylarnino, N-methylcarbamoyl, N,N-dimethylcarbamoyl, methylthio, methylsulphinyl, mesyl, N-methylsulphamoyl, N,N-dimethylsulphamoyl, N-methylsulphamoylamino and N,N-dimethylsulphamoylamino;

\mathbf{R}^{20} , \mathbf{R}^{24} , \mathbf{R}^{26} , \mathbf{R}^{30} or \mathbf{R}^{35} are independently selected from $\text{C}_{1-6}\text{alkyl}$, $\text{C}_{1-6}\text{alkanoyl}$, $\text{C}_{1-6}\text{alkylsulphonyl}$, $\text{C}_{1-6}\text{alkoxycarbonyl}$, carbamoyl, $N-(\text{C}_{1-6}\text{alkyl})\text{carbamoyl}$, $N,N-(\text{C}_{1-6}\text{alkyl})\text{carbamoyl}$, benzyl, benzyloxycarbonyl, benzoyl and phenylsulphonyl; or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

- 5 Further suitable compounds possessing IBAT inhibitory activity have the following structure of formula (CI):



(CI)

wherein:

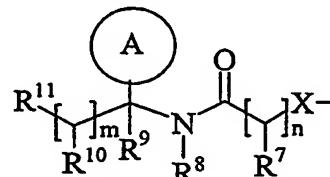
- 10 One of \mathbf{R}^1 and \mathbf{R}^2 are selected from hydrogen or $\text{C}_{1-6}\text{alkyl}$ and the other is selected from $\text{C}_{1-6}\text{alkyl}$;

\mathbf{R}^y is selected from hydrogen, hydroxy, $\text{C}_{1-6}\text{alkyl}$, $\text{C}_{1-4}\text{alkoxy}$ and $\text{C}_{1-6}\text{alkanoyloxy}$;

- 15 \mathbf{R}^z is selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, $\text{C}_{1-6}\text{alkyl}$, $\text{C}_{2-6}\text{alkenyl}$, $\text{C}_{2-6}\text{alkynyl}$, $\text{C}_{1-6}\text{alkoxy}$, $\text{C}_{1-6}\text{alkanoyl}$, $\text{C}_{1-6}\text{alkanoyloxy}$, $N-(\text{C}_{1-6}\text{alkyl})\text{amino}$, $N,N-(\text{C}_{1-6}\text{alkyl})_2\text{amino}$, $\text{C}_{1-6}\text{alkanoylamino}$, $N-(\text{C}_{1-6}\text{alkyl})\text{carbamoyl}$, $N,N-(\text{C}_{1-6}\text{alkyl})_2\text{carbamoyl}$, $\text{C}_{1-6}\text{alkylS(O)}_a$ wherein a is 0 to 2, $\text{C}_{1-6}\text{alkoxycarbonyl}$, $N-(\text{C}_{1-6}\text{alkyl})\text{sulphamoyl}$ and $N,N-(\text{C}_{1-6}\text{alkyl})_2\text{sulphamoyl}$;

v is 0-5;

one of \mathbf{R}^4 and \mathbf{R}^5 is a group of formula (CIA):



(CIA)

20

\mathbf{R}^3 and \mathbf{R}^6 and the other of \mathbf{R}^4 and \mathbf{R}^5 are independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, $\text{C}_{1-4}\text{alkyl}$, $\text{C}_{2-4}\text{alkenyl}$, $\text{C}_{2-4}\text{alkynyl}$, $\text{C}_{1-4}\text{alkoxy}$, $\text{C}_{1-4}\text{alkanoyl}$, $\text{C}_{1-4}\text{alkanoyloxy}$, $N-(\text{C}_{1-4}\text{alkyl})\text{amino}$,

N,N-(C₁₋₄alkyl)₂amino, C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl,

N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a wherein a is 0 to 2, C₁₋₄alkoxycarbonyl,

N-(C₁₋₄alkyl)sulphamoyl and N,N-(C₁₋₄alkyl)₂sulphamoyl; wherein R³ and R⁶ and the other of R⁴ and R⁵ may be optionally substituted on carbon by one or more R¹⁶;

5 X is -O-, -N(R^a)-, -S(O)_b- or -CH(R^a)-; wherein R^a is hydrogen or C₁₋₆alkyl and b is 0-2;

Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from R¹⁷;

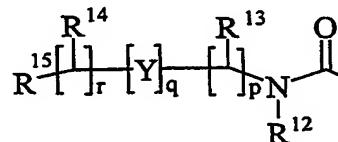
10 R⁷ is hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein R⁷ is optionally substituted by one or more substituents selected from R¹⁸;

R⁸ is hydrogen or C₁₋₄alkyl;

R⁹ is hydrogen or C₁₋₄alkyl;

R¹⁰ is hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein R¹⁰ is optionally substituted by one or more substituents selected from R¹⁹;

15 R¹¹ is carboxy, sulpho, sulphino, phosphono, -P(O)(OR^c)(OR^d), -P(O)(OH)(OR^c), -P(O)(OH)(R^d) or -P(O)(OR^c)(R^d) wherein R^c and R^d are independently selected from C₁₋₆alkyl; or R¹¹ is a group of formula (CIB):



(CIB)

20 wherein:

Y is -N(R^x)-, -N(R^x)C(O)-, -O-, and -S(O)a-; wherein a is 0-2 and R^x is hydrogen or C₁₋₄alkyl;

R¹² is hydrogen or C₁₋₄alkyl;

25 R¹³ and R¹⁴ are independently selected from hydrogen, C₁₋₄alkyl, carbocyclyl or heterocyclyl; wherein R¹³ and R¹⁴ may be independently optionally substituted by one or more substituents selected from R²⁰;

R¹⁵ is carboxy, sulpho, sulphino, phosphono, -P(O)(OR^e)(OR^f), -P(O)(OH)(OR^e), -P(O)(OH)(R^e) or -P(O)(OR^e)(R^f) wherein R^e and R^f are independently selected from C₁₋₆alkyl;

30 p is 1-3; wherein the values of R¹³ may be the same or different;
q is 0-1;

r is 0-3; wherein the values of R¹⁴ may be the same or different;

m is 0-2; wherein the values of R¹⁰ may be the same or different;

n is 1-3; wherein the values of R⁷ may be the same or different;

R¹⁶, R¹⁷ and R¹⁸ are independently selected from halo, nitro, cyano, hydroxy, amino,

5 carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy,

C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino,

C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a,

wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl and

N,N-(C₁₋₄alkyl)₂sulphamoyl; wherein R¹⁶, R¹⁷ and R¹⁸ may be independently optionally

10 substituted on carbon by one or more R²¹;

R¹⁹ and R²⁰ are independently selected from halo, nitro, cyano, hydroxy, amino,

carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy,

C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino,

C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a,

15 wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl,

N,N-(C₁₋₄alkyl)₂sulphamoyl, carbocyclyl, heterocyclyl, sulpho, sulphino, amidino, phosphono,

-P(O)(OR^a)(OR^b), -P(O)(OH)(OR^a), -P(O)(OH)(R^a) or -P(O)(OR^a)(R^b), wherein R^a and R^b are

independently selected from C₁₋₆alkyl; wherein R¹⁹ and R²⁰ may be independently optionally

substituted on carbon by one or more R²²;

20 R²¹ and R²² are independently selected from halo, hydroxy, cyano, carbamoyl, ureido,

amino, nitro, carboxy, carbamoyl, mercapto, sulphamoyl, trifluoromethyl, trifluoromethoxy,

methyl, ethyl, methoxy, ethoxy, vinyl, allyl, ethynyl, methoxycarbonyl, formyl, acetyl,

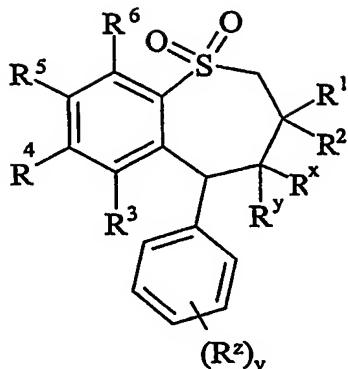
formamido, acetylamino, acetoxy, methylamino, dimethylamino, N-methylcarbamoyl,

N,N-dimethylcarbamoyl, methylthio, methylsulphinyl, mesyl, N-methylsulphamoyl and

25 N,N-dimethylsulphamoyl;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Further suitable compounds possessing IBAT inhibitory activity have the following
structure of formula (DI):



wherein:

One of R^1 and R^2 are selected from hydrogen or C_{1-6} alkyl and the other is selected

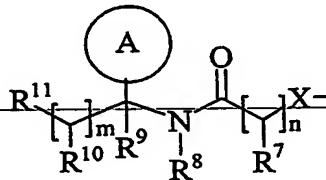
5 from C_{1-6} alkyl;

R^x and R^y are independently selected from hydrogen, hydroxy, amino, mercapto, C_{1-6} alkyl, C_{1-6} alkoxy, $N-(C_{1-6}$ alkyl)amino, $N,N-(C_{1-6}$ alkyl)₂amino, C_{1-6} alkylS(O)_a wherein a is 0 to 2;

10 R^z is selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{1-6} alkoxy, C_{1-6} alkanoyl, C_{1-6} alkanoyloxy, $N-(C_{1-6}$ alkyl)amino, $N,N-(C_{1-6}$ alkyl)₂amino, C_{1-6} alkanoylamino, $N-(C_{1-6}$ alkyl)carbamoyl, $N,N-(C_{1-6}$ alkyl)₂carbamoyl, C_{1-6} alkylS(O)_a wherein a is 0 to 2, C_{1-6} alkoxycarbonyl, $N-(C_{1-6}$ alkyl)sulphamoyl and $N,N-(C_{1-6}$ alkyl)₂sulphamoyl;

v is 0-5;

15 one of R^4 and R^5 is a group of formula (DIA):



R^3 and R^6 and the other of R^4 and R^5 are independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-4} alkyl,

20 C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, C_{1-4} alkanoyl, C_{1-4} alkanoyloxy, $N-(C_{1-4}$ alkyl)amino, $N,N-(C_{1-4}$ alkyl)₂amino, C_{1-4} alkanoylamino, $N-(C_{1-4}$ alkyl)carbamoyl,

$N,N-(C_{1-4}$ alkyl)₂carbamoyl, C_{1-4} alkylS(O)_a wherein a is 0 to 2, C_{1-4} alkoxycarbonyl, $N-(C_{1-4}$ alkyl)sulphamoyl and $N,N-(C_{1-4}$ alkyl)₂sulphamoyl; wherein R^3 and R^6 and the other of R^4 and R^5 may be optionally substituted on carbon by one or more R^{16} ;

X is $-O-$, $-N(R^a)-$, $-S(O)_b-$ or $-CH(R^a)-$; wherein R^a is hydrogen or C_{1-6} alkyl and b is 0-2;

Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from R^{17} ;

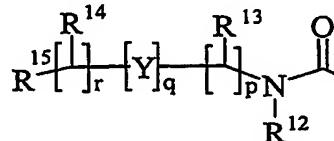
5 **R⁷** is hydrogen, C_{1-4} alkyl, carbocyclyl or heterocyclyl; wherein R^7 is optionally substituted by one or more substituents selected from R^{18} ;

R⁸ is hydrogen or C_{1-4} alkyl;

R⁹ is hydrogen or C_{1-4} alkyl;

10 **R¹⁰** is hydrogen, C_{1-4} alkyl, carbocyclyl or heterocyclyl; wherein R^{10} is optionally substituted by one or more substituents selected from R^{19} ;

R¹¹ is carboxy, sulpho, sulphino, phosphono, $-P(O)(OR^c)(OR^d)$, $-P(O)(OH)(OR^c)$, $-P(O)(OH)(R^d)$ or $-P(O)(OR^c)(R^d)$ wherein R^c and R^d are independently selected from C_{1-6} alkyl; or R^{11} is a group of formula (DIB):



15 (DIB)

wherein:

Y is $-N(R^n)-$, $-N(R^n)C(O)-$, $-O-$, and $-S(O)a-$; wherein a is 0-2 and R^n is hydrogen or C_{1-4} alkyl;

R¹² is hydrogen or C_{1-4} alkyl;

20 **R¹³** and **R¹⁴** are independently selected from hydrogen, C_{1-4} alkyl, carbocyclyl or heterocyclyl; wherein R^{13} and R^{14} may be independently optionally substituted by one or more substituents selected from R^{20} ;

R¹⁵ is carboxy, sulpho, sulphino, phosphono, $-P(O)(OR^e)(OR^f)$, $-P(O)(OH)(OR^e)$, $-P(O)(OH)(R^e)$ or $-P(O)(OR^e)(R^f)$ wherein R^e and R^f are independently selected from C_{1-6} alkyl;

25 **p** is 1-3; wherein the values of R^{13} may be the same or different;

q is 0-1;

r is 0-3; wherein the values of R^{14} may be the same or different;

m is 0-2; wherein the values of R^{10} may be the same or different;

30 **n** is 1-3; wherein the values of R^7 may be the same or different;

R^{16} , R^{17} and R^{18} are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, C_{1-4} alkanoyl, C_{1-4} alkanoyloxy, $N-(C_{1-4}\text{alkyl})\text{amino}$, $N,N-(C_{1-4}\text{alkyl})_2\text{amino}$, $C_{1-4}\text{alkanoylamino}$, $N-(C_{1-4}\text{alkyl})\text{carbamoyl}$, $N,N-(C_{1-4}\text{alkyl})_2\text{carbamoyl}$, $C_{1-4}\text{alkylS(O)}_a$

5 wherein a is 0 to 2, $C_{1-4}\text{alkoxycarbonyl}$, $N-(C_{1-4}\text{alkyl})\text{sulphamoyl}$ and $N,N-(C_{1-4}\text{alkyl})_2\text{sulphamoyl}$; wherein R^{16} , R^{17} and R^{18} may be independently optionally substituted on carbon by one or more R^{21} ;

R^{19} and R^{20} are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, C_{1-4} alkanoyl, C_{1-4} alkanoyloxy, $N-(C_{1-4}\text{alkyl})\text{amino}$, $N,N-(C_{1-4}\text{alkyl})_2\text{amino}$, $C_{1-4}\text{alkanoylamino}$, $N-(C_{1-4}\text{alkyl})\text{carbamoyl}$, $N,N-(C_{1-4}\text{alkyl})_2\text{carbamoyl}$, $C_{1-4}\text{alkylS(O)}_a$ wherein a is 0 to 2, $C_{1-4}\text{alkoxycarbonyl}$, $N-(C_{1-4}\text{alkyl})\text{sulphamoyl}$, $N,N-(C_{1-4}\text{alkyl})_2\text{sulphamoyl}$, carbocyclyl, heterocyclyl, sulpho, sulphino, amidino, phosphono, $-P(O)(OR^a)(OR^b)$, $-P(O)(OH)(OR^a)$, $-P(O)(OH)(R^a)$ or $-P(O)(OR^a)(R^b)$, wherein R^a and R^b are

10 independently selected from C_{1-6} alkyl; wherein R^{19} and R^{20} may be independently optionally substituted on carbon by one or more R^{22} ;

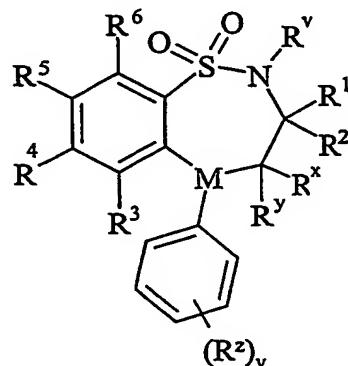
15

R^{21} and R^{22} are independently selected from halo, hydroxy, cyano, carbamoyl, ureido, amino, nitro, carboxy, carbamoyl, mercapto, sulphamoyl, trifluoromethyl, trifluoromethoxy, methyl, ethyl, methoxy, ethoxy, vinyl, allyl, ethynyl, methoxycarbonyl, formyl, acetyl, 20 formamido, acetylarnino, acetoxy, methylarnino, dimethylarnino, N -methylcarbamoyl, N,N -dimethylcarbamoyl, methylthio, methylsulphinyl, mesyl, N -methylsulphamoyl and N,N -dimethylsulphamoyl;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Further suitable compounds possessing IBAT inhibitory activity have the following

25 structure of formula (EI):



wherein:

R^y is selected from hydrogen or C_{1-6} alkyl;

One of R^1 and R^2 are selected from hydrogen or C_{1-6} alkyl and the other is selected from C_{1-6} alkyl;

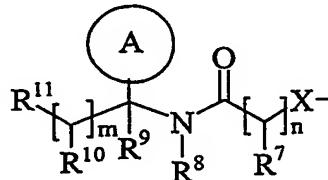
5 R^x and R^y are independently selected from hydrogen, hydroxy, amino, mercapto, C_{1-6} alkyl, C_{1-6} alkoxy, $N-(C_{1-6}$ alkyl)amino, $N,N-(C_{1-6}$ alkyl)₂amino, C_{1-6} alkylS(O)_a wherein a is 0 to 2;

 M is selected from -N- or -CH-;

10 R^z is selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{1-6} alkoxy, C_{1-6} alkanoyl, C_{1-6} alkanoyloxy, $N-(C_{1-6}$ alkyl)amino, $N,N-(C_{1-6}$ alkyl)₂amino, C_{1-6} alkanoylamino, $N-(C_{1-6}$ alkyl)carbamoyl, $N,N-(C_{1-6}$ alkyl)₂carbamoyl, C_{1-6} alkylS(O)_a wherein a is 0 to 2, C_{1-6} alkoxycarbonyl, $N-(C_{1-6}$ alkyl)sulphamoyl and $N,N-(C_{1-6}$ alkyl)₂sulphamoyl;

 v is 0-5;

15 one of R^4 and R^5 is a group of formula (EIA):



(EIA)

20 R^3 and R^6 and the other of R^4 and R^5 are independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, C_{1-4} alkanoyl, C_{1-4} alkanoyloxy, $N-(C_{1-4}$ alkyl)amino, $N,N-(C_{1-4}$ alkyl)₂amino, C_{1-4} alkanoylamino, $N-(C_{1-4}$ alkyl)carbamoyl, $N,N-(C_{1-4}$ alkyl)₂carbamoyl, C_{1-4} alkylS(O)_a wherein a is 0 to 2, C_{1-4} alkoxycarbonyl, $N-(C_{1-4}$ alkyl)sulphamoyl and $N,N-(C_{1-4}$ alkyl)₂sulphamoyl; wherein R^3 and R^6 and the other of R^4 and R^5 may be optionally substituted on carbon by one or more R^{16} ;

25 X is -O-, -N(R^a)-, -S(O)_b- or -CH(R^a)-; wherein R^a is hydrogen or C_{1-6} alkyl and b is 0-2;

 Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted by one or more substituents selected from R^{17} ;

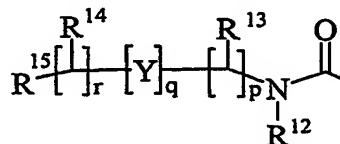
30 R^7 is hydrogen, C_{1-4} alkyl, carbocyclyl or heterocyclyl; wherein R^7 is optionally substituted by one or more substituents selected from R^{18} ;

R^8 is hydrogen or C_{1-4} alkyl;

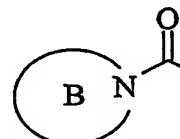
R^9 is hydrogen or C_{1-4} alkyl;

R^{10} is hydrogen, C_{1-4} alkyl, carbocyclyl or heterocyclyl; wherein R^{10} is optionally substituted by one or more substituents selected from R^{19} ;

5 R^{11} is carboxy, sulpho, sulphino, phosphono, $-P(O)(OR^c)(OR^d)$, $-P(O)(OH)(OR^c)$, $-P(O)(OH)(R^d)$ or $-P(O)(OR^c)(R^d)$ wherein R^c and R^d are independently selected from C_{1-6} alkyl; or R^{11} is a group of formula (EIB) or (EIC):



(EIB)



(EIC)

10 wherein:

Y is $-N(R^n)$, $-N(R^n)C(O)$, $-N(R^n)C(O)(CR^sR^t)$, $N(R^n)C(O)-$, $-O-$, and $-S(O)a-$;

wherein a is 0-2, v is 1-2, R^s and R^t are independently selected from hydrogen or C_{1-4} alkyl optionally substituted by R^{26} and R^n is hydrogen or C_{1-4} alkyl;

R^{12} is hydrogen or C_{1-4} alkyl;

15 R^{13} and R^{14} are independently selected from hydrogen, C_{1-4} alkyl, carbocyclyl or heterocyclyl; and when q is 0, R^{14} may additionally be selected from hydroxy; wherein R^{13} and R^{14} may be independently optionally substituted by one or more substituents selected from R^{20} ;

20 R^{15} is carboxy, sulpho, sulphino, phosphono, $-P(O)(OR^e)(OR^f)$, $-P(O)(OH)(OR^e)$, $-P(O)(OH)(R^e)$ or $-P(O)(OR^e)(R^f)$ wherein R^e and R^f are independently selected from C_{1-6} alkyl;

p is 1-3; wherein the values of R^{13} may be the same or different;

q is 0-1;

r is 0-3; wherein the values of R^{14} may be the same or different;

25 m is 0-2; wherein the values of R^{10} may be the same or different;

n is 1-3; wherein the values of R^7 may be the same or different;

Ring B is a nitrogen linked heterocyclyl substituted on carbon by one group selected from R^{23} , and optionally additionally substituted on carbon by one or more R^{24} ; and wherein if said nitrogen linked heterocyclyl contains an $-NH-$ moiety, that nitrogen may be optionally substituted by a group selected from R^{25} ;

30

R^{16} , R^{17} and R^{18} are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino, C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a 5 wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl and N,N-(C₁₋₄alkyl)₂sulphamoyl; wherein R¹⁶, R¹⁷ and R¹⁸ may be independently optionally substituted on carbon by one or more R²¹;

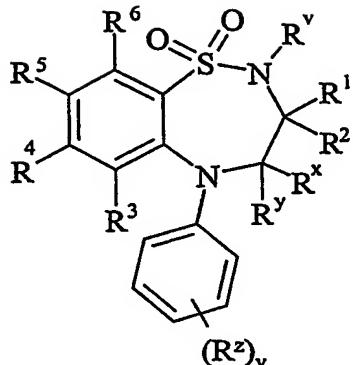
10 R^{19} , R^{20} , R^{24} and R^{26} are independently selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, N-(C₁₋₄alkyl)amino, N,N-(C₁₋₄alkyl)₂amino, C₁₋₄alkanoylamino, N-(C₁₋₄alkyl)carbamoyl, N,N-(C₁₋₄alkyl)₂carbamoyl, C₁₋₄alkylS(O)_a wherein a is 0 to 2, C₁₋₄alkoxycarbonyl, N-(C₁₋₄alkyl)sulphamoyl, N,N-(C₁₋₄alkyl)₂sulphamoyl, carbocyclyl, heterocyclyl, benzyloxycarbonylamino, sulpho, sulphino, amidino, phosphono, -P(O)(OR^a)(OR^b), -P(O)(OH)(OR^a), -P(O)(OH)(R^a) or 15 -P(O)(OR^a)(R^b), wherein R^a and R^b are independently selected from C₁₋₆alkyl; wherein R¹⁹, R²⁰, R²⁴ and R²⁶ may be independently optionally substituted on carbon by one or more R²²;

20 R^{21} and R^{22} are independently selected from halo, hydroxy, cyano, carbamoyl, ureido, amino, nitro, carboxy, carbamoyl, mercapto, sulphamoyl, trifluoromethyl, trifluoromethoxy, methyl, ethyl, methoxy, ethoxy, vinyl, allyl, ethynyl, methoxycarbonyl, formyl, acetyl, formamido, acetylarnino, acetoxy, methylarnino, dimethylarnino, N-methylcarbamoyl, N,N-dimethylcarbamoyl, methylthio, methylsulphinyl, mesyl, N-methylsulphamoyl and N,N-dimethylsulphamoyl;

25 R^{23} is carboxy, sulpho, sulphino, phosphono, -P(O)(OR^g)(OR^h), -P(O)(OH)(OR^g), -P(O)(OH)(R^g) or -P(O)(OR^g)(R^h) wherein R^g and R^h are independently selected from C₁₋₆alkyl;

30 R^{25} is selected from C₁₋₆alkyl, C₁₋₆alkanoyl, C₁₋₆alkylsulphonyl, C₁₋₆alkoxycarbonyl, carbamoyl, N-(C₁₋₆alkyl)carbamoyl, N,N-(C₁₋₆alkyl)carbamoyl, benzyl, benzyloxycarbonyl, benzoyl and phenylsulphonyl; or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

Further suitable compounds possessing IBAT inhibitory activity have the following structure of formula (FFI):



(FI)

wherein:

R^v is selected from hydrogen or C₁₋₆alkyl;

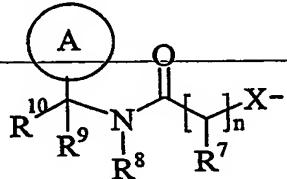
5 One of **R¹** and **R²** are selected from hydrogen or C₁₋₆alkyl and the other is selected from C₁₋₆alkyl;

R^x and **R^y** are independently selected from hydrogen, hydroxy, amino, mercapto, C₁₋₆alkyl, C₁₋₆alkoxy, *N*-(C₁₋₆alkyl)amino, *N,N*-(C₁₋₆alkyl)₂amino, C₁₋₆alkylS(O)_a wherein a is 0 to 2;

10 **R^z** is selected from halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₁₋₆alkoxy, C₁₋₆alkanoyl, C₁₋₆alkanoyloxy, *N*-(C₁₋₆alkyl)amino, *N,N*-(C₁₋₆alkyl)₂amino, C₁₋₆alkanoylamino, *N*-(C₁₋₆alkyl)carbamoyl, *N,N*-(C₁₋₆alkyl)₂carbamoyl, C₁₋₆alkylS(O)_a wherein a is 0 to 2, C₁₋₆alkoxycarbonyl, *N*-(C₁₋₆alkyl)sulphamoyl and *N,N*-(C₁₋₆alkyl)₂sulphamoyl;

15 **v** is 0-5;

one of **R⁴** and **R⁵** is a group of formula (FIA):



(FIA)

R³ and **R⁶** and the other of **R⁴** and **R⁵** are independently selected from hydrogen, halo, 20 nitro, cyano, hydroxy, amino, carboxy, carbamoyl, mercapto, sulphamoyl, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₁₋₆alkoxy, C₁₋₆alkanoyl, C₁₋₆alkanoyloxy, *N*-(C₁₋₆alkyl)amino, *N,N*-(C₁₋₆alkyl)₂amino, C₁₋₆alkanoylamino, *N*-(C₁₋₆alkyl)carbamoyl, *N,N*-(C₁₋₆alkyl)₂carbamoyl, C₁₋₆alkylS(O)_a wherein a is 0 to 2, C₁₋₆alkoxycarbonyl,

N-(C₁₋₆alkyl)sulphamoyl and *N,N*-(C₁₋₆alkyl)₂sulphamoyl; wherein R³ and R⁶ and the other of R⁴ and R⁵ may be optionally substituted on carbon by one or more R¹⁷;

X is -O-, -N(R^a)-, -S(O)_b- or -CH(R^a)-; wherein R^a is hydrogen or C₁₋₆alkyl and b is 0-2;

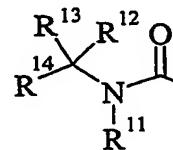
5 Ring A is aryl or heteroaryl; wherein Ring A is optionally substituted on carbon by one or more substituents selected from R¹⁸;

R⁷ is hydrogen, C₁₋₆alkyl, carbocyclyl or heterocyclyl; wherein R⁷ is optionally substituted on carbon by one or more substituents selected from R¹⁹; and wherein if said heterocyclyl contains an -NH- group, that nitrogen may be optionally substituted by a group selected from R²⁰;

R⁸ is hydrogen or C₁₋₆alkyl;

R⁹ is hydrogen or C₁₋₆alkyl;

R¹⁰ is hydrogen, halo, nitro, cyano, hydroxy, amino, carbamoyl, mercapto, sulphamoyl, hydroxyaminocarbonyl, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₁₋₁₀alkoxy, C₁₋₁₀alkanoyl, C₁₋₁₀alkanoyloxy, *N*-(C₁₋₁₀alkyl)amino, *N,N*-(C₁₋₁₀alkyl)₂amino, 15 *N,N,N*-(C₁₋₁₀alkyl)₃ammonio, C₁₋₁₀alkanoylamino, *N*-(C₁₋₁₀alkyl)carbamoyl, *N,N*-(C₁₋₁₀alkyl)₂carbamoyl, C₁₋₁₀alkylS(O)_a wherein a is 0 to 2, *N*-(C₁₋₁₀alkyl)sulphamoyl, *N,N*-(C₁₋₁₀alkyl)₂sulphamoyl, *N*-(C₁₋₁₀alkyl)sulphamoylamino, *N,N*-(C₁₋₁₀alkyl)₂sulphamoylamino, C₁₋₁₀alkoxycarbonylamino, carbocyclyl, 20 carbocyclylC₁₋₁₀alkyl, heterocyclyl, heterocyclylC₁₋₁₀alkyl, carbocyclyl-(C₁₋₁₀alkylene)_p-R²¹-(C₁₋₁₀alkylene)_q- or heterocyclyl-(C₁₋₁₀alkylene)_r-R²²-(C₁₋₁₀alkylene)_s-; wherein R¹⁰ is optionally substituted on carbon by one or more substituents selected from R²³; and wherein if said heterocyclyl contains an -NH- group, that nitrogen may be optionally substituted by a group selected from R²⁴; or R¹⁰ is a group of formula (FIB):



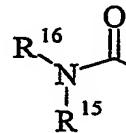
(FIB)

wherein:

R¹¹ is hydrogen or C₁₋₆alkyl;

30 R¹² and R¹³ are independently selected from hydrogen, halo, carbamoyl, sulphamoyl, C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₁₋₁₀alkanoyl, *N*-(C₁₋₁₀alkyl)carbamoyl,

- N,N-(C₁₋₁₀alkyl)₂carbamoyl, C₁₋₁₀alkylS(O)_a* wherein *a* is 0 to 2, *N-(C₁₋₁₀alkyl)sulphamoyl,*
N,N-(C₁₋₁₀alkyl)₂sulphamoyl, N-(C₁₋₁₀alkyl)sulphamoylamino,
N,N-(C₁₋₁₀alkyl)₂sulphamoylamino, carbocyclyl or heterocyclyl; wherein R¹² and R¹³ may be
independently optionally substituted on carbon by one or more substituents selected from R²⁵;
5 *and wherein if said heterocyclyl contains an -NH- group, that nitrogen may be optionally*
substituted by a group selected from R²⁶;
- R¹⁴ is selected from hydrogen, halo, carbamoyl, sulphamoyl, hydroxyaminocarbonyl,
C₁₋₁₀alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₁₋₁₀alkanoyl, *N-(C₁₋₁₀alkyl)carbamoyl,*
N,N-(C₁₋₁₀alkyl)₂carbamoyl, C₁₋₁₀alkylS(O)_a wherein *a* is 0 to 2, *N-(C₁₋₁₀alkyl)sulphamoyl,*
10 *N,N-(C₁₋₁₀alkyl)₂sulphamoyl, N-(C₁₋₁₀alkyl)sulphamoylamino,*
N,N-(C₁₋₁₀alkyl)₂sulphamoylamino, carbocyclyl, carbocyclylC₁₋₁₀alkyl, heterocyclyl,
heterocyclylC₁₋₁₀alkyl, carbocyclyl-(C₁₋₁₀alkylene)_p-R²⁷-(C₁₋₁₀alkylene)_q- or
heterocyclyl-(C₁₋₁₀alkylene)_r-R²⁸-(C₁₋₁₀alkylene)_s-; wherein R¹⁴ may be optionally substituted
15 *on carbon by one or more substituents selected from R²⁹; and wherein if said heterocyclyl*
contains an -NH- group, that nitrogen may be optionally substituted by a group selected from
R³⁰; or R¹⁴ is a group of formula (FIC):



(FIC)

- R¹⁵ is hydrogen or C₁₋₆alkyl;
20 R¹⁶ is hydrogen or C₁₋₆alkyl; wherein R¹⁶ may be optionally substituted on carbon by
one or more groups selected from R³¹;

n is 1-3; wherein the values of R⁷ may be the same or different;

- R¹⁷, R¹⁸, R¹⁹, R²³, R²⁵, R²⁹ or R³¹ are independently selected from halo, nitro, cyano,
hydroxy, amino, carbamoyl, mercapto, sulphamoyl, hydroxyaminocarbonyl, C₁₋₁₀alkyl,
25 C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, C₁₋₁₀alkoxy, C₁₋₁₀alkanoyl, C₁₋₁₀alkanoyloxy, *N-(C₁₋₁₀alkyl)amino,*
N,N-(C₁₋₁₀alkyl)₂amino, N,N,N-(C₁₋₁₀alkyl)₃ammonio, C₁₋₁₀alkanoylamino,
N-(C₁₋₁₀alkyl)carbamoyl, N,N-(C₁₋₁₀alkyl)₂carbamoyl, C₁₋₁₀alkylS(O)_a wherein *a* is 0 to 2,
N-(C₁₋₁₀alkyl)sulphamoyl, N,N-(C₁₋₁₀alkyl)₂sulphamoyl, N-(C₁₋₁₀alkyl)sulphamoylamino,
N,N-(C₁₋₁₀alkyl)₂sulphamoylamino, C₁₋₁₀alkoxycarbonylamino, carbocyclyl,
30 *carbocyclylC₁₋₁₀alkyl, heterocyclyl, heterocyclylC₁₋₁₀alkyl,*
carbocyclyl-(C₁₋₁₀alkylene)_p-R³²-(C₁₋₁₀alkylene)_q- or

heterocyclyl-(C₁₋₁₀alkylene)-R³³-(C₁₋₁₀alkylene)_s-; wherein R¹⁷, R¹⁸, R¹⁹, R²³, R²⁵, R²⁹ or R³¹ may be independently optionally substituted on carbon by one or more R³⁴; and wherein if said heterocyclyl contains an -NH- group, that nitrogen may be optionally substituted by a group selected from R³⁵;

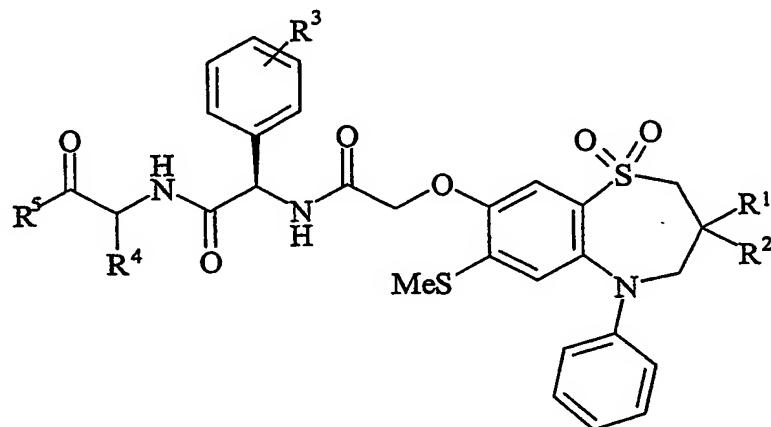
5 R²¹, R²², R²⁷, R²⁸, R³² or R³³ are independently selected from -O-, -NR³⁶-, -S(O)_x-, -NR³⁶C(O)NR³⁶-, -NR³⁶C(S)NR³⁶-, -OC(O)N=C-, -NR³⁶C(O)- or -C(O)NR³⁶-, wherein R³⁶ is selected from hydrogen or C₁₋₆alkyl, and x is 0-2;

 p, q, r and s are independently selected from 0-2;

10 R³⁴ is selected from halo, hydroxy, cyano, carbamoyl, ureido, amino, nitro, carbamoyl, mercapto, sulphamoyl, trifluoromethyl, trifluoromethoxy, methyl, ethyl, methoxy, ethoxy, vinyl, allyl, ethynyl, formyl, acetyl, formamido, acetylarnino, acetoxy, methylarnino, dimethylarnino, N-methylcarbamoyl, N,N-dimethylcarbamoyl, methylthio, methylsulphinyl, mesyl, N-methylsulphamoyl, N,N-dimethylsulphamoyl, N-methylsulphamoylarnino and N,N-dimethylsulphamoylarnino;

15 R²⁰, R²⁴, R²⁶, R³⁰ or R³⁵ are independently selected from C₁₋₆alkyl, C₁₋₆alkanoyl, C₁₋₆alkylsulphonyl, C₁₋₆alkoxycarbonyl, carbamoyl, N-(C₁₋₆alkyl)carbamoyl, N,N-(C₁₋₆alkyl)carbamoyl, benzyl, benzyloxycarbonyl, benzoyl and phenylsulphonyl; or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

A compound of formula (GI):



20

(GI)

wherein:

R¹ and R² are independently selected from C₁₋₄alkyl;

R³ is hydrogen, hydroxy or halo;

R^4 is C_{1-4} alkyl optionally substituted by hydroxy, methoxy and methylS(O)a wherein a is 0-2

R^5 is hydroxy or $HOC(O)CH(R^6)NH-$;

R^6 is selected from hydrogen and C_{1-3} alkyl optionally substituted by hydroxy,

- 5 methoxy and methylS(O)a wherein a is 0-2;
or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof;
with the proviso that when R^1 and R^2 are both butyl, R^5 is hydroxy and R^4 is
methylthiomethyl, methylsulphinylmethyl, methylthiomethyl, hydroxymethyl,
methoxymethyl; R^3 is not hydrogen; and with the proviso that when R^1 and R^2 are both butyl,
10 R^5 is $HOC(O)CH(R^6)NH-$, R^6 is hydroxymethyl and R^4 is hydroxymethyl; R^3 is not hydrogen.

Compounds of formula (AI), (BI), (CI), (DI), (EI), (FI) and (GI) or a
pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof may be
prepared by processes known in the art.

- In a particular aspect of the invention an IBAT inhibitor or a pharmaceutically
15 acceptable salt, solvate, solvate of such a salt or a prodrug thereof is an IBAT inhibitor or a
pharmaceutically acceptable salt thereof.

Suitable pharmaceutically acceptable salts of the above compounds are, for example,
an acid-addition salt of a compound of the invention which is sufficiently basic, for example,
an acid-addition salt with, for example, an inorganic or organic acid, for example
20 hydrochloric, hydrobromic, sulphuric, phosphoric, trifluoroacetic, citric, acetate or maleic
acid. In addition a suitable pharmaceutically acceptable salt of a compound which is
sufficiently acidic is an alkali metal salt, for example a sodium or potassium salt, an alkaline
earth metal salt, for example a calcium or magnesium salt, an ammonium salt or a salt with an
organic base which affords a physiologically-acceptable cation, for example a salt with
25 methylamine, dimethylamine, trimethylamine, piperidine, morpholine or
tris-(2-hydroxyethyl)amine.

The compounds may be administered in the form of a pro-drug which is broken down
in the human or animal body to give the parent compound. Examples of pro-drugs include *in*
vivo hydrolysable esters and *in vivo* hydrolysable amides.

- 30 An *in vivo* hydrolysable ester of a compound containing carboxy or hydroxy group is,
for example, a pharmaceutically acceptable ester which is hydrolysed in the human or animal
body to produce the parent acid or alcohol. Suitable pharmaceutically acceptable esters for
carboxy include C_{1-6} alkoxymethyl esters for example methoxymethyl, C_{1-6} alkanoyloxymethyl

esters for example pivaloyloxymethyl, phthalidyl esters, C₃₋₈cycloalkoxycarbonyloxyC₁₋₆alkyl esters for example 1-cyclohexylcarbonyloxyethyl; 1,3-dioxolen-2-onylmethyl esters for example 5-methyl-1,3-dioxolen-2-onylmethyl; and C₁₋₆alkoxycarbonyloxyethyl esters for example 1-methoxycarbonyloxyethyl and may be formed at any carboxy group in the compounds.

An *in vivo* hydrolysable ester of a compound containing a hydroxy group includes inorganic esters such as phosphate esters and α -acyloxyalkyl ethers and related compounds which as a result of the *in vivo* hydrolysis of the ester breakdown to give the parent hydroxy group. Examples of α -acyloxyalkyl ethers include acetoxymethoxy and 2,2-dimethylpropionyloxy-methoxy. A selection of *in vivo* hydrolysable ester forming groups for hydroxy include alkanoyl, benzoyl, phenylacetyl and substituted benzoyl and phenylacetyl, alkoxycarbonyl (to give alkyl carbonate esters), dialkylcarbamoyl and N-(dialkylaminoethyl)-N-alkylcarbamoyl (to give carbamates), dialkylaminoacetyl and carboxyacetyl. Examples of substituents on benzoyl include morpholino and piperazino-linked from a ring nitrogen atom via a methylene group to the 3- or 4- position of the benzoyl ring.

A suitable value for an *in vivo* hydrolysable amide of a compound containing a carboxy group is, for example, a N-C₁₋₆alkyl or N,N-di-C₁₋₆alkyl amide such as N-methyl, N-ethyl, N-propyl, N,N-dimethyl, N-ethyl-N-methyl or N,N-diethyl amide.

20 **Experimental**

Colon fistulated dogs may be used to demonstrate the effectiveness of the combination of the present invention in preventing diarrhoea. The IBAT inhibitor is dosed orally at a dose that will cause diarrhoea, for example 25-50 μ mol/kg. The metal salt is then introduced into the colon, through the fistulae, to see if the diarrhoea can be prevented. The dose of the metal salt varies and can be determined after analysing the bile acid concentration in faeces from dogs having been exposed to the same dose of the IBAT inhibitor.

According to another feature of the invention there is provided the use of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in combination with a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, in the manufacture of a medicament for use in the production of an IBAT inhibitory effect in a warm-blooded animal, such as man.

Suitably the production of an IBAT inhibitory effect means the treatment of hyperlipidaemic conditions. Suitably the production of an IBAT inhibitory effect means the

treatment of dyslipidemic conditions and disorders such as hyperlipidaemia, hypertriglyceridemia, hyperbetalipoproteinemia (high LDL), hyperprebetalipoproteinemia (high VLDL), hyperchylomicronemia, hypolipoproteinemia, hypercholesterolemia, hyperlipoproteinemia and hypoalphalipoproteinemia (low HDL). Suitably the production of
5 an IBAT inhibitory effect means the treatment of different clinical conditions such as atherosclerosis, arteriosclerosis, arrhythmia, hyper-thrombotic conditions, vascular dysfunction, endothelial dysfunction, heart failure, coronary heart diseases, cardiovascular diseases, myocardial infarction, angina pectoris, peripheral vascular diseases, inflammation of cardiovascular tissues such as heart, valves, vasculature, arteries and veins, aneurisms,
10 stenosis, restenosis, vascular plaques, vascular fatty streaks, leukocyte, monocytes and/or macrophage infiltrate, intimal thickening, medial thinning, infectious and surgical trauma and vascular thrombosis, stroke and transient ischaemic attacks. Suitably the production of an IBAT inhibitory effect means the treatment of atherosclerosis, coronary heart diseases,
15 myocardial infarction, angina pectoris, peripheral vascular diseases, stroke and transient ischaemic attacks.

According to another feature of the invention there is provided the use of a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, in the manufacture of a medicament for use in preventing diarrhoea that would result from excess bile acids in the intestine following administration of an IBAT inhibitor, or a
20 pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, which medicament comprises an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in combination with a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon.

According to a further feature of this aspect of the invention there is provided a
25 method for producing an IBAT inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in combination with a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon.

30 Therefore according to the present invention, there is provided a method of preventing diarrhoea that would result from excess bile acids in the intestine following administration of an effective amount an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a warm blooded animal, such as man, in need of such

treatment, which comprises administering to said animal said effective amount of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof in combination with a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon.

5 According to a further aspect of the invention there is provided a pharmaceutical composition which comprises an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in combination with a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, in association with a pharmaceutically acceptable diluent or carrier.

10 According to a further aspect of the invention there is provided a pharmaceutical composition which comprises an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in combination with a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, in association with a pharmaceutically acceptable diluent or carrier for use in producing an 15 IBAT inhibitory effect, in a warm-blooded animal, such as man.

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in combination with a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, in 20 association with a pharmaceutically acceptable diluent or carrier for use in preventing diarrhoea that would result from excess bile acids in the intestine following administration of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a warm-blooded animal, such as man.

The pharmaceutical compositions may be in a form suitable for oral administration, 25 for example as a tablet or capsule or for rectal administration as a suppository. In general the above compositions may be prepared in a conventional manner using conventional excipients.

According to an additional feature of the invention, there is provided an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in combination with a metal salt, wherein the metal salt is formulated to release in the 30 terminal ileum, caecum and/or the colon, for use as a medicament.

According to an additional feature of the invention, there is provided an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in combination with a metal salt, wherein the metal salt is formulated to release in the

terminal ileum, caecum and/or the colon, for use in producing an IBAT inhibitory effect, in a warm-blooded animal, such as man.

According to an additional feature of the invention, there is provided an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in combination with a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, for use in preventing diarrhoea that would result from excess bile acids in the intestine following administration of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a warm-blooded animal, such as man.

According to a further aspect of the present invention there is provided a kit comprising an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, optionally with instructions for use.

According to a further aspect of the present invention there is provided a kit comprising an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, optionally with instructions for use; for use in producing an IBAT inhibitory effect, in a warm-blooded animal, such as man.

According to a further aspect of the present invention there is provided a kit comprising an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon; optionally with instructions for use; for use in preventing diarrhoea that would result from excess bile acids in the intestine following administration of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a warm-blooded animal, such as man.

According to a further aspect of the present invention there is provided a kit comprising:

- a) an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a first unit dosage form;
- b) a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon; in a second unit dosage form; and
- c) container means for containing said first and second dosage forms; and optionally
- d) with instructions for use.

According to a further aspect of the present invention there is provided a kit comprising:

a) an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a first unit dosage form;

5 b) a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon; in a second unit dosage form; and

c) container means for containing said first and second dosage forms; and optionally

d) with instructions for use;

for use in producing an IBAT inhibitory effect, in a warm-blooded animal, such as man.

10 According to a further aspect of the present invention there is provided a kit comprising:

a) an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a first unit dosage form;

b) a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon; in a second unit dosage form; and

15 c) container means for containing said first and second dosage forms; and optionally

d) with instructions for use;

for use in preventing diarrhoea that would result from excess bile acids in the intestine following administration of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a warm-blooded animal, such as man.

20 According to a further aspect of the present invention there is provided a combination comprising an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, for use in producing an IBAT inhibitory effect, in a warm-blooded animal, such as man.

25 According to a further aspect of the present invention there is provided a combination comprising an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, for use in preventing diarrhoea that would result from excess bile acids in the intestine following administration of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a warm-blooded animal, such as man.

According to a further aspect of the present invention there is provided a combination treatment comprising the administration of an effective amount of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable diluent or carrier, in combination with 5 an effective amount of a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, optionally together with a pharmaceutically acceptable diluent or carrier to a warm-blooded animal, such as man in need of such therapeutic treatment.

According to a further aspect of the present invention there is provided a combination 10 treatment comprising the administration of an effective amount of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable diluent or carrier, in combination with an effective amount of a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, optionally together with a pharmaceutically acceptable diluent or carrier to a warm-blooded animal, such as man in need of such 15 therapeutic treatment for use in producing an IBAT inhibitory effect, in a warm-blooded animal, such as man.

According to a further aspect of the present invention there is provided a combination treatment comprising the administration of an effective amount of an IBAT inhibitor, or a 20 pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, optionally together with a pharmaceutically acceptable diluent or carrier, in combination with an effective amount of a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, optionally together with a pharmaceutically acceptable diluent or carrier to a warm-blooded animal, such as man in need of such 25 therapeutic treatment for use in preventing diarrhoea that would result from excess bile acids in the intestine following administration of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a warm-blooded animal, such as man.

The IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a 30 salt or a prodrug thereof, will normally be administered to a warm-blooded animal at a unit dose within the range 5-5000 mg per square meter body area of the animal, i.e. approximately 0.01-50 mg/kg, and this would be expected to provide a therapeutically-effective dose. A unit dose from such as a tablet or capsule will usually contain, for example 1-250 mg of active

ingredient. In one aspect of the invention a daily dose in the range of 0.02-50 mg/kg is employed. In another aspect a daily dose in the range of 0.02-20 mg/kg is employed. However the daily dose will necessarily be varied depending upon the host treated, the particular route of administration, and the severity of the illness being treated. Accordingly the optimum dosage may be determined by the practitioner who is treating any particular patient.

The metal salt will normally be administered to a warm-blooded animal at a unit dose within the range of up to 500mg/kg and this would be expected to provide a therapeutically-effective dose. In one aspect of the invention a daily dose in the range of up to 100mg/kg per day is employed. However the daily dose will necessarily be varied depending upon the host treated, the particular route of administration, and the severity of the illness being treated. Accordingly the optimum dosage may be determined by the practitioner who is treating any particular patient. Suitably this dose will be 1g or less per patient per day. More suitably it will be 500mg or less per patient per day. In another aspect a daily dose in the range of 50-100 mg per day is employed.

The dosage of each of the two drugs and their proportions have to be composed so that the best possible treatment effects, as defined by national and international guidelines (which are periodically reviewed and re-defined), will be met.

The combination therapy defined hereinbefore may also involve, in addition to the combination, one or more other substances and/or treatments. Such conjoint treatment may be achieved by way of the simultaneous, sequential or separate administration of the individual components of the treatment.

Suitable additional substances include HMG Co-A reductase inhibitors, or pharmaceutically acceptable salts, solvates, solvates of such salts or prodrugs thereof. Suitable HMG Co-A reductase inhibitors, pharmaceutically acceptable salts, solvates, solvates of such salts or prodrugs thereof are statins well known in the art. Particular statins are fluvastatin, lovastatin, pravastatin, simvastatin, atorvastatin, cerivastatin, bervastatin, dalvastatin, mevastatin and rosuvastatin, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof. A particular statin is atorvastatin, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof. A more particular statin is atorvastatin calcium salt. A further particular statin is rosuvastatin, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof. A preferable particular statin is rosuvastatin calcium salt.

Further suitable additional substances include:

- a CETP (cholesteryl ester transfer protein) inhibitor, for example those referenced and described in WO 00/38725 page 7 line 22 - page 10, line 17 which are incorporated herein by reference;
- a cholesterol absorption antagonist for example azetidinones such as SCH 58235 and those described in US 5,767,115 which are incorporated herein by reference;
- a MTP (microsomal transfer protein) inhibitor for example those described in Science, 282, 751-54, 1998 which are incorporated herein by reference;
- a fibric acid derivative; for example clofibrate, gemfibrozil, fenofibrate, ciprofibrate and bezafibrate;
- a nicotinic acid derivative, for example, nicotinic acid (niacin), acipimox and nericitrol;
- a phytosterol compound for example stanols;
- probucol;
- an anti-obesity compound for example orlistat (EP 129,748) and sibutramine (GB 2,184,122 and US 4,929,629);
- an antihypertensive compound for example an angiotensin converting enzyme (ACE) inhibitor, an angiotensin II receptor antagonist, an adrenergic blocker, an alpha adrenergic blocker, a beta adrenergic blocker, a mixed alpha/beta adrenergic blocker, an adrenergic stimulant, calcium channel blocker, a diuretic or a vasodilator;
- insulin;
- sulphonylureas including glibenclamide, tolbutamide;
- metformin; and/or
- acarbose;

or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof,

optionally together with a pharmaceutically acceptable diluent or carrier to a warm-blooded animal, such as man in need of such therapeutic treatment.

Particular ACE inhibitors or pharmaceutically acceptable salts, solvates, solvate of such salts or a prodrugs thereof, including active metabolites, which can be used as an additional substance include but are not limited to, the following compounds: alacepril, alatriopril, altiopril calcium, ancovenin, benazepril, benazepril hydrochloride, benazeprilat, benzoylcaptopril, captopril, captopril-cysteine, captopril-glutathione, ceranapril, ceranopril, ceronapril, cilazapril, cilazaprilat, delapril, delapril-diacid, enalapril, enalaprilat, enapril, epicaptopril, foroxymithine, fosfenopril, fosenopril, fosenopril sodium, fosinopril, fosinopril

sodium, fosinoprilat, fosinoprilic acid, glycopril, hemorphin-4, idrapril, imidapril, indolapril, indolaprilat, libenzapril, lisinopril, lyciumin A, lyciumin B, mixanpril, moexipril, moexiprilat, moveltipril, muracein A, muracein B, muracein C, pentopril, perindopril, perindoprilat, pivalopril, pivopril, quinapril, quinapril hydrochloride, quinaprilat, ramipril, ramiprilat, spirapril, spirapril hydrochloride, spiraprilat, spiropril, spiropril hydrochloride, temocapril, temocapril hydrochloride, teprotide, trandolapril, trandolaprilat, utibapril, zabicipril, zabiciprilat, zofenopril and zofenoprilat. Preferred ACE inhibitors for use in the present invention are ramipril, ramiprilat, lisinopril, enalapril and enalaprilat. More preferred ACE inhibitors for uses in the present invention are ramipril and ramiprilat.

10 Preferred angiotensin II antagonists, pharmaceutically acceptable salts, solvates, solvate of such salts or a prodrugs thereof for use as an additional substance, but are not limited to candesartan, candesartan cilexetil, losartan, valsartan, irbesartan, tasosartan, telmisartan and eprosartan. Particularly preferred angiotensin II antagonists or pharmaceutically acceptable derivatives thereof for use in the present invention are
15 candesartan and candesartan cilexetil.

Additional suitable additional substances are PPAR alpha and/or gamma agonists, or pharmaceutically acceptable salts, solvates, solvates of such salts or prodrugs thereof. Suitable PPAR alpha and/or gamma agonists, pharmaceutically acceptable salts, solvates, solvates of such salts or prodrugs thereof are well known in the art. These include the compounds
20 described in WO 01/12187, WO 01/12612, WO 99/62870, WO 99/62872, WO 99/62871, WO 98/57941, WO 01/40170, J Med Chem, 1996, 39, 665, Expert Opinion on Therapeutic Patents, 10 (5), 623-634 (in particular the compounds described in the patent applications listed on page 634) and J Med Chem, 2000, 43, 527 which are all incorporated herein by reference. Particularly a PPAR alpha and/or gamma agonist refers to WY-14643, clofibrate, 25 fenofibrate, bezafibrate, GW 9578, troglitazone, pioglitazone, rosiglitazone, eglitazone, proglitazone, BRL-49634, KRP-297, JTT-501, SB 213068, GW 1929, GW 7845, GW 0207, L-796449, L-165041 and GW 2433. Particularly a PPAR alpha and/or gamma agonist refers to (S)-2-ethoxy-3-[4-(2-{4-methanesulphonyloxyphenyl}ethoxy)phenyl] propanoic acid and pharmaceutically acceptable salts thereof.

30 The metal salt can be formulated in a delayed release single or multiple unit oral formulation. The delayed release of the metal salt can be achieved by for example using techniques producing formulations with time dependent or pH dependent release or enzymatically degradable formulations (Pharmaceutics. The Science of Dosage Form Design

Second Edition; Ed. Micheal E Aulton; Harcourt Publishers Limited; 2002). These formulations can be manufactured with conventional techniques, for example as described in Aulton,(see above), or Industrial Aspects of Pharmaceutics, Ed Erik Sandell; Swedish Pharmaceutical Press; 1993).

- 5 The IBAT inhibitor may be formulated by conventional techniques.

Claims

1. A combination which comprises an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a metal salt, wherein the metal
5 salt is formulated to release in the terminal ileum, caecum and/or the colon.
2. A combination according to claim 1 wherein the metal salt is a calcium salt.
3. A combination according to either of claims 1 or 2 wherein the metal salt is calcium
10 phosphate.
4. A combination according to any one of claims 1 - 3 wherein the IBAT inhibitor is a benzothiepine.
- 15 5. A combination according to any one of claims 1 - 4 wherein the IBAT inhibitor is selected from:

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(carboxymethyl) carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(carboxymethyl)carbamoyl]-4-
20 hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(2-sulphoethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)-1'-phenyl-1'-[N'-(2-sulphoethyl)carbamoyl]methyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
25 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-sulphoethyl)carbamoyl]-4- hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-sulphoethyl) carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-
30 carboxyethyl)carbamoyl]benzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-carboxyethyl)carbamoyl]-4- hydroxybenzyl}carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;

- 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(5-carboxypentyl)carbamoyl]benzyl} carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-carboxyethyl)carbamoyl]benzyl} carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 5 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{\mathbf{\alpha}}-[N'-(2-sulphoethyl)carbamoyl]-2-fluorobenzyl} carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(R)-(2-hydroxy-1-carboxyethyl)carbamoyl]benzyl} carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(R)-(2-hydroxy-1-carboxyethyl)carbamoyl]benzyl} carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 10 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{N-[(R)- α -{N'-{(R)-1-[N''-(R)-(2-hydroxy-1-carboxyethyl)carbamoyl]-2-hydroxyethyl}carbamoyl}benzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{\mathbf{\alpha}}-[N'-(carboxymethyl)carbamoyl]benzyl} carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 15 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-(N-{\mathbf{\alpha}}-[N'-(ethoxy)(methyl)phosphoryl-methyl)carbamoyl]benzyl} carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3-butyl-3-ethyl-5-phenyl-7-methylthio-8-{N-[(R)- α -{N'-{2-[(hydroxy)(methyl)phosphoryl]ethyl}carbamoyl]benzyl}carbamoylmethoxy}-2,3,4,5-
- 20 tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[N'-(2-methylthio-1-carboxyethyl)carbamoyl]benzyl} carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{N-[(R)- α -{N'-{2-[(methyl)(ethyl)phosphoryl]ethyl}carbamoyl}-4-hydroxybenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-
-
- 25 benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-{N-[(R)- α -{N'-{2-[(methyl)(hydroxy)phosphoryl]ethyl}carbamoyl}-4-hydroxybenzyl]carbamoylmethoxy}-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 1,1-dioxo-3,3-dibutyl-5-phenyl-7-methylthio-8-(N-{(R)- α -[(R)-N'-(2-methylsulphinyl-1-carboxyethyl)carbamoyl]benzyl} carbamoylmethoxy)-2,3,4,5-tetrahydro-1,5-benzothiazepine;
- 30 and

1,1-dioxo-3,3-dibutyl-5-phenyl-7-methoxy-8-[N-{(R)- α -[N'-(2-sulphoethyl)carbamoyl]-4-hydroxybenzyl}carbamoylmethoxy]-2,3,4,5-tetrahydro-1,5-benzothiazepine;
or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.

- 5 6. The use of a combination according to any one of claims 1-5, in the manufacture of a medicament for use in the production of an IBAT inhibitory effect in a warm-blooded animal, such as man.
- 10 7. The use of a combination according to any one of claims 1-5, in the manufacture of a medicament for use in preventing diarrhoea that would result from excess bile acids in the intestine following administration of an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof.
- 15 8. A method for producing an IBAT inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a combination according to any one of claims 1-5.
- 20 9. A method of preventing diarrhoea that would result from excess bile acids in the intestine following administration of an effective amount an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, in a warm blooded animal, such as man, in need of such treatment, which comprises administering to said animal said effective amount of a combination according to any one of claims 1-5.
- 25 10. A pharmaceutical composition which comprises a combination according to any one of claims 1-5, in association with a pharmaceutically acceptable diluent or carrier.
11. A combination according to any one of claims 1-5 for use as a medicament.

A B S T R A C TTITLE : THERAPEUTIC TREATMENT

5 A combination comprising an IBAT inhibitor, or a pharmaceutically acceptable salt, solvate, solvate of such a salt or a prodrug thereof, and a metal salt, wherein the metal salt is formulated to release in the terminal ileum, caecum and/or the colon, is described. Compositions containing this combination and uses of the combination are also described.

PATENT COOPERATION TREATY

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From the
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

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**NOTIFICATION OF TRANSMITTAL OF
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Applicant

ASTRAZENECA AB et al.

1. The applicant is hereby notified that this International Preliminary Examining Authority transmits herewith the international preliminary examination report and its annexes, if any, established on the international application.
2. A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.
3. Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report (but not of any annexes) and will transmit such translation to those Offices.

4. REMINDER

The applicant must enter the national phase before each elected Office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices)(Article 39(1))(see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details on the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

The applicant's attention is drawn to Article 33(5), which provides that the criteria of novelty, inventive step and industrial applicability described in Article 33(2) to (4) merely serve the purposes of international preliminary examination and that "any Contracting State may apply additional or different criteria for the purposes of deciding whether, in that State, the claimed inventions is patentable or not" (see also Article 27(5)). Such additional criteria may relate, for example, to exemptions from patentability, requirements for enabling disclosure, clarity and support for the claims.

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